1 1. A composition providing sustained release of a drug, the composition comprising 2 a mucopolysaccharide, a carrier protein, and a drug. 1 2. The composition of claim 1, wherein the composition consists of the mucopolysaccharide, the carrier protein, the drug, and one or more pharmaceutically 2 3 acceptable additives. 1 3. The composition of claim 1, wherein the ratio of the total mass of mucopolysaccharide in the composition to the total mass of carrier protein in the composition 2 3 is about 1:1 to 1:20. 1 4. The composition of claim 1, wherein the mucopolysaccharide is chondroitin sulfate or hyaluronate. 5. The composition of claim 1, wherein the carrier protein is a γ -globulin, albumin, fibrinogen, histone, protamine, gelatin, or collagen. 6. The composition of claim 1, wherein the carrier protein is a γ -globulin. 1 1 7. The composition of claim 1, wherein the carrier protein is an albumin. 1 8. The composition of claim 1, wherein the drug is a protein drug. 1 9. The composition of claim 8, wherein the protein drug is an erythropoietin, 2 granulocyte colony stimulating factor, granulocyte macrophage colony stimulating factor, thrombopoietin, interferon-α, interferon-β, interferon-γ, urokinase, tissue plasminogen 3 4 activator, interleukin-11, fibroblast growth factor, epidermal growth factor, growth hormone, 5 brain-derived neurotrophic factor, nerve growth factor, leptin, neurotrophin-3, superoxide 6 dismutase, antibody, calcitonin, insulin, or parathyroid hormone.

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1 2	10. The composition of claim 1, wherein the composition contains about 0.1 to 50% by weight the mucopolysaccharide.
1	11. The composition of claim 1, wherein the composition contains about 0.1 to 2%
2	by weight the drug.
1	12. A method of producing a sustained release drug composition, the method
2	comprising
3	providing a precipitating solution containing a mucopolysaccharide, a carrier protein,
4	and a drug;
5	lowering the pH of the precipitating solution to a level sufficient to form an insoluble
6	product comprising the mucopolysaccharide, the carrier protein, and the drug; and
7	collecting from the precipitating solution the insoluble product.
1	13. The method of claim 12, wherein the insoluble product consists of the
2	mucopolysaccharide, the carrier protein, the drug, and one or more pharmaceutically
3	acceptable additives.
1	14. The method of claim 12, wherein the ratio of the total mass of
2	mucopolysaccharide in the insoluble product to the total mass of carrier protein in the
3	insoluble product is about 1:1 to 1:20.
1	15. The method of claim 12, wherein the mucopolysaccharide is chondroitin sulfate
2	or hyaluronate.
1	16. The method of claim 12, wherein the carrier protein is a γ-globulin, albumin,
2	fibrinogen, histone, protamine, gelatin, or collagen.
1	17. The method of claim 12, wherein the carrier protein is a γ -globulin.
1	18. The method of claim 12, wherein the carrier protein is an albumin

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1	19. The inethod of claim 12, wherein the drug is a protein drug.
1	20. The method of claim 12, wherein the protein drug is an erythropoietin,
2	granulocyte colony stimulating factor, granulocyte-macrophage colony stimulating factor,
3	thrombopoietin, interferon-α, interferon-β, interferon-γ, urokinase, tissue plasminogen
4	activator, interleukin-11, fibroblast growth factor, epidermal growth factor, growth hormone,
5	brain-derived neurotrophic factor, nerve growth factor, leptin, neurotrophin-3, superoxide
6	dismutase, antibody, calcitonin, insulin, or parathyroid hormone.
1	21. The method of claim 12, wherein the pH of the solution is about 7 or above
2	before the lowering step.
1	22. The method of claim 12, wherein the pH of the solution is lowered to about 2 to 4
2	in the lowering step.
1	23. The method of claim 12, further comprising, prior to the providing step, mixing a
2	first solution containing the carrier protein and the drug with a second solution containing the
3	mucopolysaccharide to produce the precipitating solution.
1	24. The method of claim 12, wherein the precipitating solution contains zinc or
2	calcium ions.
1	25. The method of claim 12, further comprising
2	suspending the insoluble product in a preparatory solution having a pH of about 6 to 8
3	to form a mixture; and
4	lyophilizing the mixture to obtain a solid product.
1	26. A composition providing sustained release of a drug, the composition comprising
2	a mucopolysaccharide and a protein drug.

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27. The composition of claim 26, wherein the composition consists of the
mucopolysaccharide, the protein drug, and one or more pharmaceutically acceptable
additives.

- 28. The composition of claim 26, wherein the mucopolysaccharide is chondroitin sulfate or hyaluronate.
- 29. The composition of claim 26, wherein the protein drug is an erythropoietin, granulocyte colony stimulating factor, granulocyte-macrophage colony stimulating factor, thrombopoietin, interferon- α , interferon- β , interferon- γ , urokinase, tissue plasminogen activator, interleukin-11, fibroblast growth factor, epidermal growth factor, growth hormone, brain-derived neurotrophic factor, nerve growth factor, leptin, neurotrophin-3, superoxide dismutase, antibody, calcitonin, insulin, or parathyroid hormone.
- 30. The composition of claim 26, wherein the composition contains about 0.1 to 50% by weight the mucopolysaccharide.
- 31. The composition of claim 26, wherein the composition contains about 0.1 to 50% by weight the protein drug.
- 32. A method of producing a sustained release drug composition, the method comprising
 - providing a precipitating solution containing a mucopolysaccharide and a protein drug;
- lowering the pH of the precipitating solution to a level sufficient to form an insoluble product comprising the mucopolysaccharide and the protein drug; and collecting from the precipitating solution the insoluble product.
- 33. The method of claim 32, wherein the insoluble product consists of the mucopolysaccharide, the protein drug, and one or more pharmaceutically acceptable additives.

1	34. The method of claim 32, wherein the mucopolysaccharide is chondroitin sulfate
2	or hyaluronate.
1	35. The method of claim 32, wherein the protein drug is an erythropoietin,
2	granulocyte colony stimulating factor, granulocyte-macrophage colony stimulating factor,
3	thrombopoietin, interferon- α , interferon- β , interferon- γ , urokinase, tissue plasminogen
4	activator, interleukin-11, fibroblast growth factor, epidermal growth factor, growth hormone,
5	brain-derived neurotrophic factor, nerve growth factor, leptin, neurotrophin-3, superoxide
6	dismutase, antibody, calcitonin, insulin, or parathyroid hormone.
1	36. The method of claim 32, wherein the pH of the solution is about 7 or above
2	before the lowering step.
1	37. The method of claim 32, wherein the pH of the solution is lowered to about 2 to 4
2	in the lowering step.
1	38. The method of claim 32, further comprising, prior to the providing step, mixing a
2	first solution containing the protein drug with a second solution containing the
3	mucopolysaccharide to produce the precipitating solution.
1	39. The method of claim 32, wherein the precipitating solution contains zinc or
2	calcium ions.
1	40. The method of claim 32, wherein the insoluble product contains about 0.1 to 50%
2	by weight the mucopolysaccharide.
1	41. The method of claim 32, wherein the insoluble product contains about 0.1 to 50%
2	by weight the protein drug.
l	42. The method of claim 32 further comprising

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suspending the insoluble product in a preparatory solution having a pH of about 6 to 8
to form a mixture; and
lyophilizing the mixture to obtain a solid product.
43. A method of delivering a drug to a subject, the method comprising introducing
the composition of claim 1 into the subject.
44. The method of claim 43, wherein the composition is introduced subcutaneously
or intramuscularly into the subject.
45. A method of delivering a drug to a subject, the method comprising introducing
the composition of claim 26 into the subject.
46. The method of claim 45, wherein the composition is introduced subcutaneously
or intramuscularly into the subject.

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